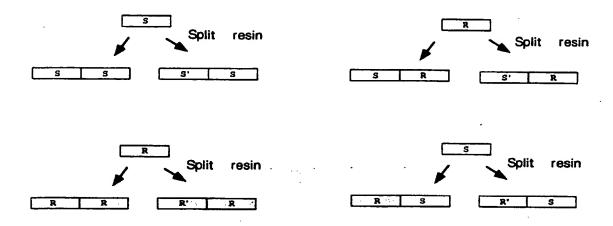
# FIGURE 1

# Generating Molecular Diversity using NCL\*

CXC chemokine S S S S SDF1 $\alpha$ 

### 8x N-terminal modules



### 4x C-terminal modules

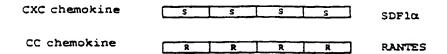


Ligation at Xxx-Cys bond to generate hybrid molecule \*NCL=native chemical ligation

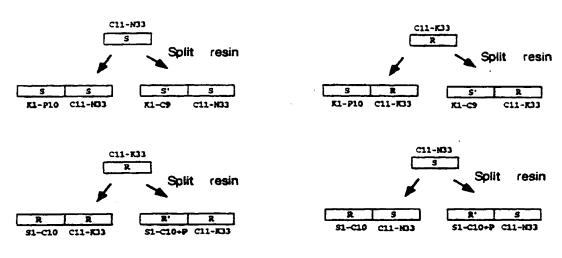
S'=-Pro & R=+Pro

FIGURE 2

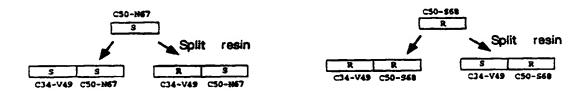
# Generating Molecular Diversity using NCL\*



# 8x N-terminal modules



### 4x C-terminal modules



Ligation at Xxx-Cys bond to generate hybrid molecule \*NCL=native chemical ligation

S'=-Pro & R'=+Pro

FIGURE 3

# CHEMOKINE PATTERNS AND MPBV

The two amino acids preceding the central cysteine are evaluated when designing improved agonists or antagonists. MPBV\* (vMIP-I or vMIP-II)

	_	>	1	*
	H	z	۸۲	
	A	Ą	õ	
20	~	æ		
	ď	Α	Ь	
	L	^	٦.	*
	Ь	Н	Ь	
	2	S	R	
	٧	E	K	
	_	F	0	<b>←</b>
	7	F	Y	
	FA	R	57	
	၁	၁	၁	
01	၁	Ь	၁	
	Ь	ပ	ᅩ	
	T	æ	D	
	T	Y	Ь	
	Q	S	~	
	S	IJ	H	
	S	S	*	
	>	>	S	
	Ъ	d	A	
	S	×	Ŋ	
	RANTES	SDF1	MPBV* G	

1	<b></b>	<b>,</b> ,			
		Ж	×	ĸ	
			z		
		24	×	×	
		ī	L	T	*
		>	R	T	
	40	궈	Ą	F	*
-		>	^	I	
		>	1	Λ	*
		Ą	0	G	
		۵		Ь	
		z	L	Ж	<b>+</b>
		S	A	S	
		ပ	၁	С	
		×	z	L	<b>←</b>
		g	Ь	δ	<b>←</b>
		S	Т	S	
	30		z		
		T	L	T	
		Y	_	Ь	
		74	ᅩ	Y	
		>	н	≱	
		Э	L	S	
	24	×	X	S	
		RANTES K E	SDF1	MPBV*	

	<del></del> -			
			8	
			A	
	S		I	
<i>L</i> 9	Σ	z	٨	
	Э	Γ	Ь	
	ı	Α	Т	
	S	×	0	
	z	E	0	
	_	Г	M	
	>	Y	L	
	Э	Э	Х	
	<b>×</b>	0	K	
	>	_	^	*
	≱	≽	*	*
	7	¥	Ω	
	ᅺ	رر	¥	
	ш	×	S	
	а	Ы	×	
	z	Ω	D	
	A	П	4	*
	ပ	၁	၁	
	>	>	>	*
	O	0	0	
	~	×	×	
	z	z	g	
	RANTES N	SDF1	MPBV*	

\* = Hydrophobic core side chains, highly conserved.

Bolded positions indicate conservation between all 3 or (MPBV and another).

 $\uparrow$  . = Unique position, MPBV matches neither RANTES nor SDF1 $\alpha$ .

All three N-terminii are unique. Likewise the two positions before the central cysteine are unique.

FIGURE 4

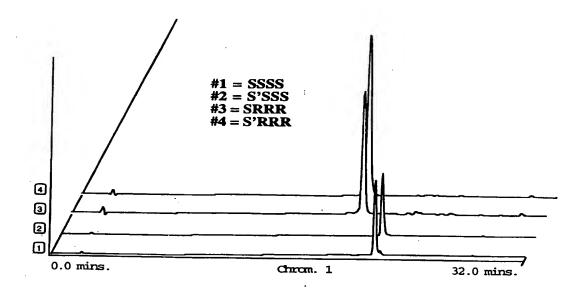


FIGURE 5

